



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

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Dossier requirements for centrally authorised products (CAPs)

Submission of applications to the European Medicines Agency, members of the Committee for Medicinal Products for Human use (CHMP), Pharmacovigilance Risk Assessment Committee (PRAC) and Committee for Advanced Therapies (CAT)*

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Application / Submission Type	Dossier Requirements for EMA, (Co-)Rapporteurs and Members/Alternates
<p>Full application</p> <p>Extension</p> <p>Renewal</p> <p>Type IB variation (with PRAC involvement). Grouping or Worksharing containing NAPs¹</p> <p>Type II variation (with or without PRAC involvement) Grouping or Worksharing containing NAPs¹</p> <p>Periodic Safety Update Report (PSUR)²</p> <p>Post Authorisation Safety Studies (PASS)</p> <p>Post-Authorisation Measures (PAMs)</p> <p>Annual Re-Assessment</p> <p>Article 20 procedure</p>	<p>All eCTD format submission for Centrally Authorised products sent to EMA via eSubmission Gateway/Web Client will be considered delivered to all National Competent Authorities' representatives, alternates and scientific experts.</p> <p>Do not submit any additional copies of eCTD format CAP submissions directly to the NCAs on CD/DVD or via CESP as this might lead to validation issues and cause delays.</p> <p>All Centralised Procedure submissions should be made via EMA eSubmission Gateway/Web Client only.</p> <p>Committee for Advanced Therapies (CAT) members nominated by the European Commission <u>not</u> linked to EMA Common Repository:</p> <p>1 copy of eCTD format submission on hard media, e.g.CD/DVD, after EMA technical validation (or by start of the content/regulatory validation phase) and after Validation Supplementary Information.</p> <p>CAT* submissions are only required if the product is an advanced therapy (AT).</p> <p>The Names and Dossier delivery addresses for CAT members nominated by the European Commission can be found here.</p>
<p>Type IA or IB variation</p> <p>Transfer</p> <p>Art 61(3) Notification</p> <p>Art. 58 (WHO) submissions</p> <p>Active Substance Master File (ASMF)</p> <p>Plasma Master File (PMF)</p>	<p>As above</p> <p>Note: No CAT* or PRAC submissions required for these procedure types.</p>

¹ Nationally authorised products concerned by this procedure should follow the dossier requirements as detailed [here](#).

**Application /
Submission Type**

Dossier Requirements for EMA, (Co-)Rapporteurs and Members/Alternates

For any other procedure type, such as Referrals submissions, PASS 107, workshare, Signal Detection and Ancillary Medical device submissions please review the

[Dossier requirements for Referrals, ASMFs and Nationally authorised products.](#)

*** for CAT, the dossier is only required by its members if the product is an advanced therapy (AT). members if the product is an advanced therapy (AT).**